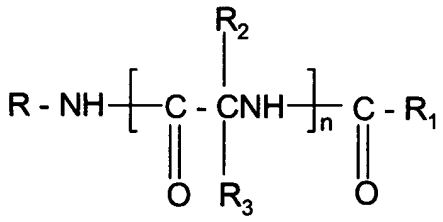
IN THE CLAIMS:

This listing of the claims replaces all prior versions and listings of the claims in the application. Please amend the Claims as indicated hereinbelow. Any claim cancelled is cancelled without prejudice.

1-19. (Cancelled)

20. (Currently Amended) A method for the prophylaxis or treatment of migraine headaches in a ~~subject,~~ patient comprising administering to said patient a headache relieving effective amount of a compound of the formula:



wherein

R is ~~hydrogen, lower alkyl, lower alkenyl, lower alkynyl, aryl,~~ aryl lower alkyl, ~~heterocyclic, heterocyclic lower alkyl, lower alkyl heterocyclic, lower cycloalkyl, lower cycloalkyl lower alkyl,~~ and R is unsubstituted or is substituted with at least one electron withdrawing group or electron donating group;

R_1 is ~~hydrogen or~~ lower alkyl[[;]], and R_1 is unsubstituted or substituted with an electron donating group or electron withdrawing group;

R_2 is ~~and R_3 are independently~~ hydrogen, lower alkyl, lower alkenyl, lower alkynyl, aryl lower alkyl, aryl, halo, heterocyclic, heterocyclic lower alkyl, lower alkyl heterocyclic, lower cycloalkyl, lower cycloalkyl lower alkyl, or $Z\text{-}Y_s$

R_3 is lower alkyl, lower alkenyl, lower alkynyl, aryl, aryl lower alkyl, halo, heterocyclic, heterocyclic lower alkyl, lower alkyl heterocyclic, lower cycloalkyl, lower cycloalkyl lower alkyl or ZY; wherein R_2 and R_3 may be unsubstituted or substituted with at least one electron withdrawing group or electron donating group and wherein heterocyclic in R_2 and R_3 is furyl, thienyl, pyrazolyl, pyrrolyl, imidazolyl, indoyl, thiazolyl, oxazolyl, isothiazolyl, isoxazolyl; piperidyl, pyrrolinyl, piperazinyl, quinolyl, triazoyl, tetrazolyl, isoquinolyl, benzofuryl, benzothienyl, morpholinyl, benzoxazolyl, tetrahydrofuryl, pyranyl, indazolyl, purinyl, indolinyl, pyrazolindinyl, imidazolinyl, imidazolindinyl, pyrrolidinyl, furazanyl, N-methylindolyl, methylfuryl, pyridazinyl, pyrimidinyl, pyrazinyl, epoxy, aziridino, oxetanyl, or azetidinyl;;

Z is O, ~~S, $S(O)_a$, NR_4, or PR_4~~ or NR_6';

Y is hydrogen, lower alkyl, aryl, aryl lower alkyl, lower alkenyl, or lower alkynyl, ~~heterocyclic, heterocyclic lower alkyl,~~ and Y may be unsubstituted or substituted with an electron donating group or an electron withdrawing group, or

ZY taken together is $NR_4NR_5R_7$, NR_4OR_5, or ONR_4R_7, ~~OPR_4R_5, PR_4OR_5, SNR_4R_7, NR_4SR_7, SPR_4R_5, or PR_4SR_7, $NR_4PR_5R_6$ or $PR_4NR_5R_7$,~~

~~$NR_4C(=O)R_5$, SCR_5, $NR_4C(=O)OR_5$, or $SC(=O)OR_5$;~~

R_4, and R_5 ~~and R_6~~ are independently hydrogen, lower alkyl, aryl, aryl lower alkyl, lower alkenyl, or lower alkynyl, wherein R_4, and R_5 ~~and R_6 may be~~ are independently unsubstituted or substituted with an electron withdrawing group or an electron donating group; and

R_6' is hydrogen or lower alkyl and R_6' may be unsubstituted or substituted with an electron withdrawing group or an electron donating group;

R_7 is $COOR_8$, COR_8, hydrogen, lower alkyl, aryl, or aryl lower alkyl~~, lower alkenyl or lower alkynyl~~, which R_7 may be unsubstituted or substituted with an electron withdrawing group or electron donating group;

R_8 is hydrogen or lower alkyl, or aryl lower alkyl, and the aryl or alkyl group may be unsubstituted or substituted with an electron withdrawing group or an electron donating group; and

~~n is 1-4;~~ n is 1;

~~a is 1-3;~~

wherein

~~heterocyclic contains from 3 up to 18 ring atoms and up to a total of 17 ring carbon atoms containing 1 to 4 hetero ring atoms selected from the group consisting of nitrogen, oxygen and sulfur.~~ the electron withdrawing group and electron donating group are selected from the group consisting of halo, nitro, lower alkenyl, lower alkynyl, formyl, aryl, trifluoromethyl, aryl lower alkanoyl, hydroxy, lower alkoxy, lower alkyl, mercapto, lower alkyl-thio and lower alkyldithio.

21. (Currently Amended) The method according to Claim 20 wherein ~~one of~~ R_2 ~~and R_3~~ is hydrogen.

22-24. (Cancelled)

25. (Currently Amended) The method according to Claim 20 wherein R_2 ~~and R_3 are~~ ~~independently~~ is hydrogen, lower alkyl, aryl, aryl lower alkyl, heterocyclic, heterocyclic lower alkyl or ZY; and

R_3 is lower alkyl, aryl, aryl lower alkyl, heterocyclic, heterocyclic lower alkyl or ZY; wherein R_2 and R_3 are independently unsubstituted or substituted by an electron withdrawing group or electron donating group. ~~Z is O, NR_4 or PR_4;~~

~~Y is hydrogen, lower alkyl, aryl, aryl lower alkyl loweralkyl, heterocyclic or heterocyclic lower alkyl; or~~

~~ZY taken together is $NR_4NR_5R_7$, NR_4OR_5, ONR_4R_7, $NR_4C(=O)R_5$, or $NR_4C(=O)OR_5$; and~~

~~R_4, R_5 and R_7 are independently hydrogen, lower alkyl, aryl or aryl lower alkyl.~~

26. (Currently Amended) The method according to Claim 25 wherein R_2 is hydrogen and R_3 is lower alkyl, aryl, aryl lower alkyl, heterocyclic, heterocyclic lower alkyl, or ZY;

~~Z is O, NR_4 or PR_4;~~

~~Y is hydrogen, lower alkyl, aryl, aryl lower alkyl loweralkyl, heterocyclic or heterocyclic lower alkyl; or~~

~~ZY taken together is $NR_4R_5R_7$, NR_4OR_5, ONR_4R_7, $NR_4C(=O)R_5$, or $NR_4C(=O)OR_5$; and~~

~~R_4, R_5 and R_7 are independently hydrogen, lower alkyl, aryl or aryl lower alkyl.~~ which R_3 may be unsubstituted or substituted with an electron withdrawing group or electron donating group.

27. (Currently Amended) The method according to Claim 26 wherein

R_2 is hydrogen and R_3 is lower alkyl, which may be unsubstituted or substituted with an electron donating or electron withdrawing group~~, NR_5OR_6, or ONR_5R_7~~.

28. (Currently Amended) The method according to Claim 26 wherein R_3 is lower alkyl which is unsubstituted or substituted with hydroxy or lower alkoxy or NR_4OR_5 ~~or ONR_4R_7~~, wherein R_4, and R_5 ~~and R_7~~ are independently hydrogen or lower alkyl, R is aryl lower alkyl, which aryl group may be unsubstituted or substituted with an electron withdrawing group and R_1 is lower alkyl.

29. (Original) The method according to Claim 26 wherein R_3 is heterocyclic.

30. (Original) The method according to Claim 29 wherein heterocyclic is heteroaromatic.

31. (Original) The method according to Claim 30 wherein R_3 is furyl, pyridyl, thienyl or thiazolyl.

32. (Original) The method according to Claim 28 wherein aryl is phenyl.

33. (Original) The method according to Claim 28 wherein aryl is phenyl and is unsubstituted or substituted with halo.

34. (Currently Amended) The method according to Claim 20 wherein the compound is

(R)-N-Benzyl-2 acetamido-3-methoxy- propionamide;

O-methyl-N-acetyl-D-serine-m-fluorobenzylamide;

O-methyl-N-acetyl-D-serine-p-fluorobenzylamide;

N-acetyl-D-phenylglycinebenzylamide;

D-1,2-(N, O-dimethylhydroxylamino)-2 acetamid[[o]]e acetic acid benzylamide;

or

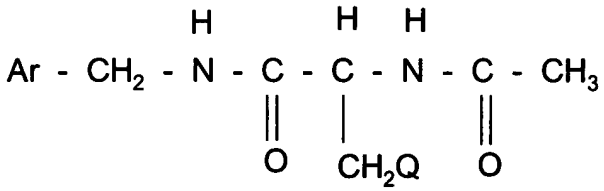
D-1,2-(O-methylhydroxylamino)-2-acetamid[[o]]e acetic acid benzylamide.

35-55. (Cancelled)

56. (Original) The method according to Claim 20 wherein the carbon atom which is substituted by R_2 and R_3 is in the D configuration.

57-62. (Cancelled)

63. (Currently Amended) The method according to Claim 20 wherein Ar is unsubstituted aryl or aryl substituted with an electron donating or electron withdrawing group and ~~halo~~ wherein the compound has the formula:



and Q is lower alkoxy.

64. (Original) The method according to Claim 63 wherein Q is methoxy.

65. (Original) The method according to Claim 63 wherein Q is methoxy and Ar is unsubstituted aryl or aryl substituted with halo.

66. (Original) The method according to Claim 63 wherein the carbon atom which is bonded to CH_2Q is in the D configuration.

67. (Original) The method according to Claim 63 wherein the carbon atom which is bonded to CH_2Q is in the D configuration.

68-72. (Cancelled)

73. (New) The method according to Claim 63 wherein Ar is unsubstituted aryl or aryl substituted with halo.

74. (New) The method according to Claim 20 wherein R is benzyl which may be unsubstituted or substituted with an electron withdrawing group or electron donating group.

75. (New) The method according to Claim 20 where R_1 is methyl.

76. (New) The method according to Claim 20 wherein R is benzyl, R_1 is lower alkyl and R_2 is hydrogen.

77. (New) The method according to Claim 76 wherein R_3 is CH_2Q, NR_4OR_5 or $NR_4NR_5R_7$, wherein Q is lower alkoxy, R_4 is hydrogen or alkyl containing 1-3 carbon atoms, R_5 is hydrogen or alkyl containing 1-3 carbon atoms and R_7 is hydrogen or alkyl containing 1-3 carbon atoms.

78. (New) The method according to Claim 77 wherein R_3 is CH_2Q.

79. (New) The method according to Claim 20 wherein R_1 is methyl, R is benzyl, R_2 is hydrogen, and R_3 is CH_2Q wherein Q is methoxy.

80. (New) The method according to Claim 20 wherein R_1 is methyl, R is m-fluorobenzyl, R_2 is H and R_3 is CH_2Q, wherein Q is methoxy.

81. (Previously Presented) The method according to Claim 20 wherein R_1 is methyl, R is p-fluorobenzyl, R_2 is H, and R_3 is CH_2Q wherein Q is methoxy.

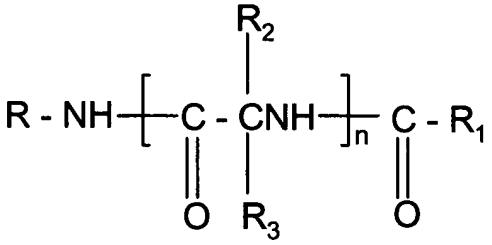
82. (New) The method according to Claim 20 wherein R_1 is methyl, R is benzyl, R_2 is hydrogen and R_3 is phenyl.

83. (New) The method according to Claim 20 wherein R_1 is methyl, R is benzyl, R_2 is hydrogen and R_3 is $N(CH_3)OCH_3$.

84. (New) The method according to Claim 20 wherein R_1 is methyl, R is benzyl , R_2 is hydrogen and R_3 is $NH(OCH_3)$.

85. (New) The method according to Claim 20 wherein R_1 is methyl, R is fluorophenyl, R_2 is H, and R_3 is CH_2Q, wherein Q is methoxy.

86. (New) A method for the prophylaxis or treatment of migraine headaches in a patient comprising administering to said patient a headache relieving effective amount of a compound of the formula:



wherein

R is aryl lower alkyl and R is unsubstituted or is substituted with at least one electron withdrawing group or electron donating group selected from the group consisting of halo, nitro, lower alkenyl, lower alkynyl, formyl, aryl, trifluoromethyl, aryl lower alkanoyl, hydroxy, lower alkoxy, lower alkyl, mercapto, lower alkylthio, and lower alkyldithio;

R_1 is methyl, and is unsubstituted or substituted with an electron donating group or an electron withdrawing group selected from the group consisting of halo, nitro, lower alkenyl,

lower alkynyl, formyl, aryl, trifluoromethyl, lower alkoxy carbonyl, aryl lower alkanoyl, hydroxy, lower alkoxy, lower alkyl, mercapto, lower alkylthio, and lower alkyldithio;

R_2 is hydrogen, lower alkyl, lower alkenyl, lower alkynyl, aryl, aryl lower alkyl, halo, heterocyclic, heterocyclic lower alkyl, lower alkyl heterocyclic, lower cycloalkyl, lower cycloalkyl lower alkyl, or ZY;

R_3 is lower alkyl, lower alkenyl, lower alkynyl, aryl, aryl lower alkyl, halo, heterocyclic, heterocyclic lower alkyl, lower alkyl heterocyclic, lower cycloalkyl, lower cycloalkyl lower alkyl or ZY;

wherein R_2 and R_3 may be unsubstituted or substituted with at least one electron withdrawing group or electron donating group and wherein heterocyclic in R_2 and R_3 is furyl, thienyl, pyrazolyl, pyrrolyl, imidazolyl, indolyl, thiazolyl, oxazolyl, isothiazolyl, isoxazolyl, piperidyl, pyrrolinyl, piperazinyl, quinolyl, triazolyl, tetrazolyl, isoquinolyl, benzofuryl, benzothienyl, morpholinyl, benzoxazolyl, tetrahydrofuryl, pyranyl, indazolyl, purinyl, indolinyl, pyrazolindinyl, imidazolinyl, imidazolindinyl, pyrrolidinyl, furazanyl, N-methylindolyl, methylfuryl, pyridazinyl, pyrimidinyl, pyrazinyl, epoxy, aziridino, oxetanyl or azetidinyl;

Z is O, or NR_6';

Y is hydrogen, lower alkyl, aryl, aryl lower alkyl, lower alkenyl or lower alkynyl, and Y may be unsubstituted or substituted with an electron donating group or an electron withdrawing group, or

ZY taken together is $NR_4NR_5R_7$, NR_4OR_5, or ONR_4R_7;

R_6' is hydrogen or lower alkyl;

R_4 and R_5 are independently hydrogen, lower alkyl, aryl, aryl lower alkyl, lower alkenyl, or lower alkynyl, and R_4 and R_5 may be independently unsubstituted or substituted with an electron withdrawing group or an electron donating group;

R_7 is $COOR_8$, COR_8, hydrogen, lower alkyl, aryl or aryl lower alkyl, which R_7 may be unsubstituted or substituted with an electron withdrawing group or electron donating group;

R_8 is hydrogen or lower alkyl, or aryl lower alkyl, and the aryl or alkyl group may be unsubstituted or substituted with an electron withdrawing group or an electron donating group; and

n is 1.

87. (New) The method according to Claim 86 wherein R_1 is methyl which is unsubstituted.

88. (New) The method according to Claim 86 wherein R is benzyl, which is unsubstituted or substituted on the phenyl ring with an electron donating group or electron withdrawing group.

89. (New) The method according to Claim 87 wherein R is benzyl, which is unsubstituted or substituted on the phenyl ring with an electron donating group or electron withdrawing group.

90. (New) The method according to Claim 86 wherein R_2 is hydrogen.

91. (New) The method according to Claim 87 wherein R_2 is hydrogen.

92. (New) The method according to Claim 88 wherein R_2 is hydrogen.

93. (New) The method according to Claim 89 wherein R_2 is hydrogen.

94. (New) The method according to Claim 86 wherein R_3 is a lower alkyl which is unsubstituted or substituted with an electron donating group or electron withdrawing group selected from the group consisting of halo, nitro, carboxy, lower alkenyl, lower alkynyl, formyl, aryl, carboxyamido, trifluoromethyl, lower alkoxycarbonyl, aryl lower alkanoyl, hydroxy, lower alkoxy, lower alkyl, amino, lower alkylamino, dilower alkylamino, aryloxy, mercapto or lower alkylthio.

95. (New) The method according to Claim 87 wherein R_3 is a lower alkyl which is unsubstituted or substituted with an electron donating group or electron withdrawing group selected from the group consisting of halo, nitro, carboxy, lower alkenyl, lower alkynyl, formyl, aryl, carboxyamido, trifluoromethyl, lower alkoxycarbonyl, aryl lower alkanoyl, hydroxy, lower alkoxy, lower alkyl, amino, lower alkylamino, dilower alkylamino, aryloxy, mercapto or lower alkylthio.

96. (New) The method according to Claim 88 wherein R_3 is a lower alkyl which is unsubstituted or substituted with an electron donating group or electron withdrawing group selected from the group consisting of halo, nitro, carboxy, lower alkenyl, lower alkynyl, formyl, aryl, carboxyamido, trifluoromethyl, lower alkoxycarbonyl, aryl lower alkanoyl, hydroxy, lower

alkoxy, lower alkyl, amino, lower alkylamino, dilower alkylamino, aryloxy, mercapto or lower alkylthio.

97. (New) The method according to Claim 89 wherein R_3 is a lower alkyl which is unsubstituted or substituted with an electron donating group or electron withdrawing group selected from the group consisting of halo, nitro, carboxy, lower alkenyl, lower alkynyl, formyl, aryl, carboxyamido, trifluoromethyl, lower alkoxycarbonyl, aryl lower alkanoyl, hydroxy, lower alkoxy, lower alkyl, amino, lower alkylamino, dilower alkylamino, aryloxy, mercapto or lower alkylthio.

98. (New) The method according to Claim 90 wherein R_3 is a lower alkyl which is unsubstituted or substituted with an electron donating group or electron withdrawing group selected from the group consisting of halo, nitro, carboxy, lower alkenyl, lower alkynyl, formyl, aryl, carboxyamido, trifluoromethyl, lower alkoxycarbonyl, aryl lower alkanoyl, hydroxy, lower alkoxy, lower alkyl, amino, lower alkylamino, dilower alkylamino, aryloxy, mercapto or lower alkylthio.

99. (New) The method according to Claim 91 wherein R_3 is a lower alkyl which is unsubstituted or substituted with an electron donating group or electron withdrawing group selected from the group consisting of halo, nitro, carboxy, lower alkenyl, lower alkynyl, formyl, aryl, carboxyamido, trifluoromethyl, lower alkoxycarbonyl, aryl lower alkanoyl, hydroxy, lower alkoxy, lower alkyl, amino, lower alkylamino, dilower alkylamino, aryloxy, mercapto or lower alkylthio.

100. (New) The method according to Claim 92 wherein R_3 is a lower alkyl which is unsubstituted or substituted with an electron donating group or electron withdrawing group selected from the group consisting of halo, nitro, carboxy, lower alkenyl, lower alkynyl, formyl, aryl, carboxyamido, trifluoromethyl, lower alkoxycarbonyl, aryl lower alkanoyl, hydroxy, lower alkoxy, lower alkyl, amino, lower alkylamino, dilower alkylamino, aryloxy, mercapto or lower alkylthio.

101. (New) The method according to Claim 93 wherein R_3 is a lower alkyl which is unsubstituted or substituted with an electron donating group or electron withdrawing group selected from the group consisting of halo, nitro, carboxy, lower alkenyl, lower alkynyl, formyl, aryl, carboxyamido, trifluoromethyl, lower alkoxycarbonyl, aryl lower alkanoyl, hydroxy, lower alkoxy, lower alkyl, amino, lower alkylamino, dilower alkylamino, aryloxy, mercapto or lower alkylthio.

102. (New) The method according to any one of Claims 86-101 wherein R_3 is lower alkyl substituted by an electron donating group.

103. (New) The method according to Claim 102 wherein R_3 is lower alkyl substituted by lower alkoxy.